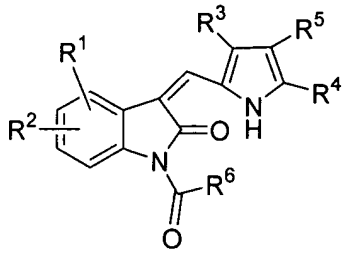
Amendments to the Claims:

This listing of claims will replace all prior versions and listings of claims in the application:

Listing of Claims:

1. (Currently amended) A compound of Formula (I):



(I)

wherein:

R^1 and R^2 are independently selected from the group consisting of hydrogen, halo, alkyl, alkylthio, nitro, trihalomethyl, hydroxy, hydroxyalkyl, alkoxy, cyano, aryl, ~~heteroaryl,~~ $-C(O)R^7$ (where R^7 is selected from the group consisting of alkyl, amino, hydroxy, alkoxy, aryl, ~~heteroaryl,~~ aryloxy, ~~heteroaryloxy, heterocycle,~~ and aminoalkylamino), $-NR^8R^9$, $-NR^8C(O)R^9$, $-SO_2R^8$, and $-S(O)_2NR^8R^9$ (where R^8 and R^9 are independently selected from the group consisting of hydrogen, alkyl and aryl~~, aryl and heteroaryl, or R^8 and R^9 together with the nitrogen to which they are attached form a saturated heterocycloamino~~);

R^3 is selected from the group consisting of hydrogen, alkyl, hydroxyalkyl, aminoalkyl, $-C(O)R^7$~~, (where R^7 is as defined above)~~, and aryl~~, aryl, and heteroaryl~~;

R^4 is selected from the group consisting of hydrogen, alkyl, $-C(O)R^7$~~, (where R^7 is as defined above)~~, and aryl~~, aryl, and heteroaryl~~;

R^5 is 3-amino-2-hydroxypropylaminocarbonyl, N-(2-dimethylaminoethyl)-aminocarbonyl, N-(2-diethylaminoethyl)-N-methylaminocarbonyl, N-(3-dimethylaminopropyl)aminocarbonyl, N-(2-diethylaminoethyl)-aminocarbonyl, N-(3-ethylaminopropyl)aminocarbonyl, N-(3-ethylamino-2-hydroxypropyl)aminocarbonyl, N-(3-diethylamino-propyl)aminocarbonyl, 3-amino-2-hydroxypropylaminocarbonyl, 3-dimethylamino-2-hydroxypropylaminocarbonyl, 3-diethylamino-2-hydroxypropylaminocarbonyl, N-(3-diethylamino-2-hydroxy-propyl)aminocarbonyl, N-(2-diethylaminoethyl)-aminocarbonyl or N-(ethylaminoethyl)aminocarbonyl;

R^6 is:

(a) $-OR^{13}$ wherein R^{13} is alkyl, trifluoromethyl, carboxyalkyl, aminoalkyl, phosphonooxyalkyl, sulfooxyalkyl, hydroxyalkyl, alkoxyalkyl, aryl, pyrrole, pyrrolidone, imidazole, thiophene, furan, tetrahydropyranyl and ~~heteroaryl, heteroaralkyl, heterocyclyl,~~ monosaccharides ~~and heterocyclylalkyl~~ wherein the alkyl chain in carboxyalkyl, aminoalkyl, phosphonooxyalkyl, sulfooxyalkyl, ~~heteroaralkyl, heterocyclylalkyl,~~ hydroxyalkyl, or alkoxyalkyl is optionally substituted with one or two hydroxy group(s) and further wherein one or two carbon atoms in said alkyl chain are optionally replaced by oxygen, $-NR^{14}-$ (where R^{14} is hydrogen or alkyl), –S-, or $-SO_2-$; or

(b) $-NR^{15}R^{16}$ where are R^{15} and R^{16} are independently selected from the group consisting of hydrogen, alkyl, carboxyalkyl, alkoxyalkyl, aminoalkyl, phosphonooxyalkyl, sulfooxyalkyl, hydroxyalkyl, pyrrole, pyrrolidone, imidazole, thiophene, furan, tetrahydropyranyl and aryl; ~~aryl, heteroaryl, heteroaralkyl, and heterocyclylalkyl;~~ wherein the alkyl chain in carboxyalkyl, aminoalkyl, phosphonooxyalkyl, ~~heteroaralkyl, heterocyclylalkyl,~~ hydroxyalkyl, or alkoxyalkyl is optionally substituted with one or two hydroxy group(s) and further wherein one or two carbon atoms in the alkyl chain are optionally replaced by oxygen, $-NR^{17}-$, ~~(where R^{17} is hydrogen or alkyl)~~, –S-, or $-SO_2-$; or R^{15} and R^{16} together with the nitrogen atom to which they are attached form a heterocycloamino, wherein the heterocycloamino is pyrrole, pyrrolidone, imidazole, thiophere or furan ~~saturated or unsaturated heterocycloamino;~~ or a pharmaceutically acceptable salt thereof.

2. (original) A pharmaceutical composition, comprising a compound or salt of Claim 1 and a pharmaceutically acceptable carrier or excipient.

3. (original) A method for the modulation of the catalytic activity of a protein kinase comprising contacting said protein kinase with a compound or salt of Claim 1.

4. (original) The method of Claim 3 wherein said protein kinase is selected from the group consisting of a receptor tyrosine kinase, a non-receptor tyrosine kinase and a serine-threonine kinase.

Claims 5-9 (canceled).

10. (original) The compound of Claim 1, wherein R^5 is N-(2-diethylaminoethyl)-aminocarbonyl.

11. (original) The compound of Claim 10, wherein R^3 and R^4 are lower alkyl having 1 to 4 carbon atoms.

12. (original) The compound of Claim 11, wherein R^3 and R^4 are methyl.

13. (original) The compound of Claim 12, wherein R^1 is hydrogen and R^2 is fluoro.

14. (original) The compound of Claim 13, wherein R^2 is a fluoro at the 5 position of the indolinone moeity.

15. (original) A pharmaceutical composition comprising the compound of Claim 14 and a pharmaceutically acceptable carrier or excipient.

Claims 16-34 (canceled)